

DESCRIPTION

PHOTOINITIATOR, NOVEL COMPOUND, AND PHOTOCURABLE COMPOSITION

5

TECHNICAL FIELD

The present invention relates to photoinitiators, a novel compound, and photocurable compositions including a photoinitiator. More specifically, the present invention relates to photoinitiators which are usable in thick-layer UV curable coatings.

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BACKGROUND ART

In recent years, photoinitiators are gaining in importance in the field of UV-light and daylight curable resin compositions, such as photocurable coatings or photocurable printing inks, which may be applicable to a wide range of substrates including metal, paper, plastics and wood, since the photoinitiators can drastically increase curing rate.

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In this field, highly sensitive photoinitiators are desired, which are capable of absorbing enough UV or daylight, and producing radicals that start polymerization or transferring the absorbed energy to polymerizable substances for formation of radicals.

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Conventionally, photoinitiators have been repeatedly studied so far with the aim of enhancing sensitivity, and improving inherent technical problems such as yellowing. In order to overcome these general problems, for example,

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alkylphenylbisacrylphosphine oxides, and photoinitiator mixtures of an alkylphenylbisacrylphosphine oxide with benzophenone are disclosed in U.S. Patent No. 6,020,528.

The conventional photoinitiators, such as
5 alkylphenylbisacrylphosphine oxides exhibit a good curability, particularly when used in thin layer photocurable coatings. However, the photoinitiators are difficult to use for UV curing of coatings having a greater layer thickness. This is because the self-absorption of the photoinitiator molecule
10 containing aromatic structure elements is large.

Furthermore, the photoinitiators of the alkylphenylbisacrylphosphine oxide type exhibit an improved yellowing behavior in comparison with other conventional photoinitiators (e.g., benzophenone). However, the problem of
15 yellowing has not been completely solved thus far. Therefore, there is a strong need for a photoinitiator showing significant improvement regarding yellowing.

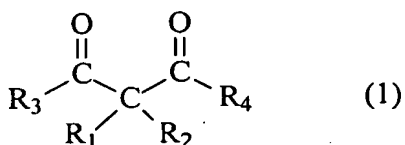
DISCLOSURE OF INVENTION

20 It is therefore an object of the invention to provide a photoinitiator which exhibits excellent photosensitivity, yields colorless products, and is also usable in thick layer UV curable coatings.

Another object is to provide a novel chemical compound
25 which is usable for the photoinitiator. Furthermore, another object is to provide a photocurable composition having aforesaid properties.

Unexpectedly, the inventors have now found that a compound having a chemical structure represented by the following formula (1) has excellent photoactivity with UV irradiation, and is well suited for photopolymerization of radical curable ethylenic unsaturated compounds.

Accordingly, the present invention provides a photoinitiator consisting essentially of a compound having a molecular weight of 1000 or less, and having a chemical structure represented by the following formula (1),



wherein R_3 and R_4 independently denote an alkyl group having a carbon number of 1 to 8, and

R_1 and R_2 independently denote

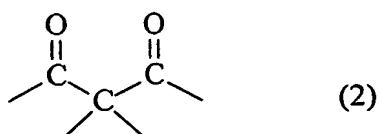
1) an electron attracting group,

2) an alkyl group having a carbon number of 1 to 8, or

3) an alkyl group having a carbon number of 1 to 8, which has an electron attracting group at the β , γ , or δ position with respect to both of the carbonyl groups, wherein the alkyl group 2) is methyl or ethyl group when each of the two

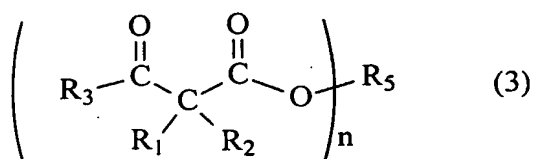
substituents is the alkyl group 2), and

mass percentage of a chemical structure element represented by the following formula (2),



which is expressed in formula (1), based on the total molecular weight of the compound having the chemical structure represented by formula (1), is within the range of 17% to 54%.

The present invention also provides a photoinitiator
 5 consisting essentially of a compound having a molecular weight of 1000 or less, and having a chemical structure represented by the following formula (3),

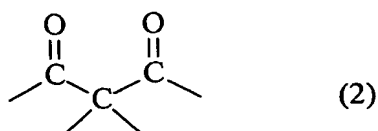


wherein R₃ denotes an alkyl group having a carbon number
 10 of 1 to 8, R₅ denotes a mono-, di-, tri-, tetra- or pentavalent aliphatic hydrocarbon group or an alkyleneoxy group-substituted aliphatic hydrocarbon group, n is a natural number of 1 to 5, and

R₁ and R₂ independently denote

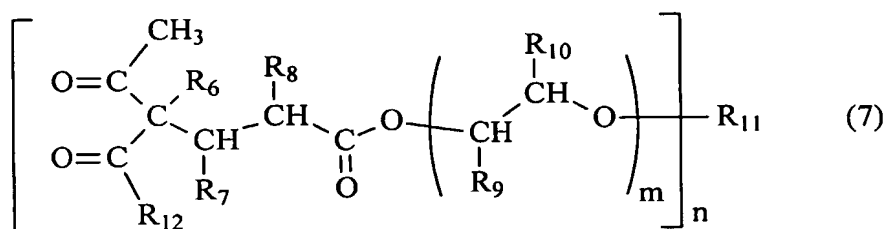
- 15 1) an electron attracting group,
- 2) an alkyl group having a carbon number of 1 to 8, or
- 3) an alkyl group having a carbon number of 1 to 8, which has an electron attracting group at the β, γ, or δ position with respect to both of the carbonyl groups, wherein the alkyl
- 20 group 2) is methyl or ethyl group when each of the two substituents is the alkyl group 2), and

weight percentage of a chemical structure element represented by the following formula (2),



which is expressed in formula (3), based on the total molecular weight of the compound having the chemical structure represented by formula (3), is within the range of 17% to 47%.

5 The present invention also provides a novel compound having a chemical structure represented by the following formula (7),



wherein

10 R_6 denotes an alkyl group having a carbon number of 1 to 8, a C_{1-4} alkyl carbonyl group, a cyano group, a C_{1-4} alkyl carbonyl methyl group, a C_{1-4} alkyl carbonyl ethyl group, a C_{1-4} alkoxy carbonyl methyl group, a C_{1-4} alkoxy carbonyl ethyl group, and an alkyl group having a carbon number of 1 to 8
15 which is substituted by carboxyl group or cyano group,

R_7 , R_8 , R_9 , and R_{10} independently denote a hydrogen atom, or a methyl group, and at least one of R_9 and R_{10} is a hydrogenatom,

R_{11} denotes a di-, tri- or tetra-valent aliphatic
20 hydrocarbon group having a carbon number of 2 to 12,

R_{12} denotes a methyl group, or an alkoxy group having a carbon number of 1 to 18,

n is a natural number of 2 to 4, and
m is an integer of 0 to 15.

The present invention also provides a photocurable
5 composition comprising,
an above-mentioned photoinitiator, and
a radical curable ethylenic unsaturated compound.

BEST MODE FOR CARRYING OUT THE INVENTION

10 R₃ and R₄, defined as an alkyl group having a carbon
number of 1 to 8 in the aforementioned chemical structure
represented by formula (1) or the aforementioned chemical
structure represented by formula (3), may be a linear or
branched alkyl group, which include methyl, ethyl, propyl,
15 isopropyl, n-butyl, t-butyl, pentyl, hexyl, cyclohexyl, n-
heptyl, and n-octyl groups. Among these structures, in
particular, a methyl group is preferable from the viewpoints
of photosensitivity.

R₅, defined as an aliphatic hydrocarbon, includes a
20 linear or branched alkyl group such as methyl, ethyl, propyl,
isopropyl, n-butyl, t-butyl, pentyl, hexyl, cyclo-hexyl, n-
heptyl, n-octyl, and a multi-functional aliphatic hydrocarbon
group such as ethylene, propylene, trimethylol propane
residual group, or pentaerythritol residual group.

25 R₅, defined as an alkyleneoxy group-substituted aliphatic
hydrocarbon group, includes an alkyl group obtained by a
reaction of methanol, ethanol, propanol, isopropanol, n-

butanol, t-butanol, pentanol, hexanol, cyclohexanol, n-heptanol, n-octanol, ethylene glycol, propylene glycol, neopentyl glycol, trimethylol propane, or pentaerythritol and ethylene oxide or propylene oxide.

5 R_1 and R_2 in the aforementioned chemical structure represented by formula (1) or the aforementioned chemical structure represented by formula (3) is selected from the group consisting of

- 1) an electron attracting group,
- 10 2) an alkyl group having a carbon number of 1 to 8, and
- 3) an alkyl group having a carbon number of 1 to 8, which has an electron attracting group at the β , γ , or δ position with respect to both of the carbonyl groups.

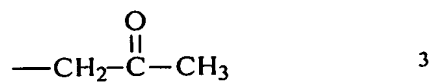
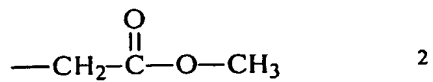
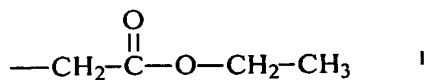
Examples of the electron attracting groups 1) include a
15 ketone group such as acetyl group, an ester group, an ether group, a carboxyl group, a cyano group, a sulfonic acid group, a sulfonyl group, or a phosphate group.

The definition of alkyl group 2), having a carbon number of 1 to 8, is the same as the definition of the above-
20 mentioned R_3 or R_4 .

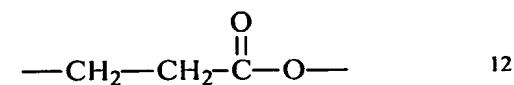
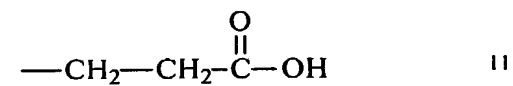
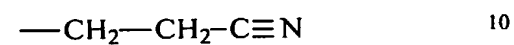
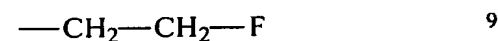
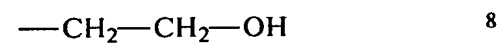
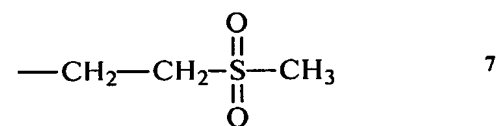
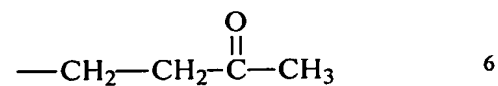
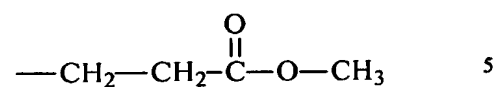
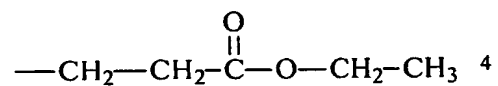
The alkyl group having a carbon number of 1 to 8, which has an electron attracting group at the β , γ , or δ position with respect to both of the carbonyl groups 3) include the following structures.

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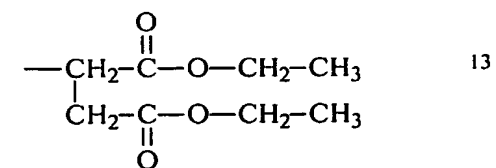
[β position substituted groups]



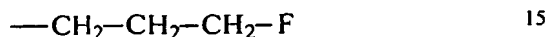
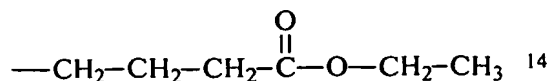
[γ position substituted groups]



[β position and γ position substituted group]



[δ position substituted groups]



Here, the γ position substituted group No. 12 may be
5 connected with another β -dicarbonyl units via an alkylene
group, a poly(alkyleneoxy) alkylene group, or a residual
structure of poly-functional archol.

In this invention, these groups may be selected randomly
as R_1 or R_2 ; however, when each of the two substituents is the
10 alkyl group 2), methyl or ethyl group must be selected as the
alkyl group 2) in order to produce at least one radical by
photoirradiation.

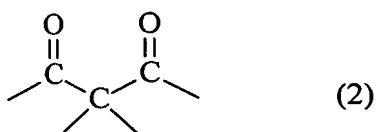
Among these above-mentioned groups, particularly
preferred groups are Nos. 4, 5 and 12, from the viewpoint of
15 photosensitivity as well as their ease of production.

In this invention, the compound having the chemical
structure represented by formula (1) or the aforementioned
chemical structure represented by formula (3), is
characterized in having a molecular weight of 1000 or less,
20 that is, a compound having the molecular weight range is
usable as an additive type photoinitiator, and furthermore,
exhibits good compatibility with radical curable monomers,
oligomers, or polymers. Furthermore, a compound having a
molecular weight in the above range can give its cured

products hardness and solvent resistance.

The preferable range of the molecular weight is 700 or less, from the viewpoint of the compatibility, and also from the viewpoint of hardness and solvent resistance of a cured
5 composition which contains the compound.

Furthermore, the compound having the chemical structure represented by formula (1) or the chemical structure represented by formula (3), is characterized in that the mass percentage of a chemical structure element represented by the
10 following formula (2),

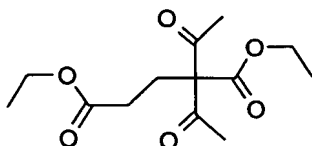


which is expressed in formula (1) or formula (3), based on the total molecular weight of the compound, is within the range of 17% to 54% by mass. The excellent photosensitivity of the
15 present invention results from the above range of the mass percentage of the chemical structure element represented by formula (2).

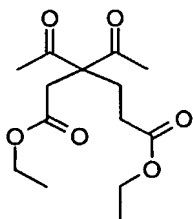
Examples of the compound having the chemical structure represented by formula (1) are

- 20 3,3-Dimethyl-2,4-pentandione,
3,3-Diethyl-2,4-pentandione,
3-Acetyl-3-methyl-4-oxo-pentanoic acid ethyl ester,
4-Acetyl-4-methyl-5-oxo-hexanoic acid ethyl ester,
4-Acetyl-4-methyl-5-oxo-hexanoic acid butyl ester,
25 4-Acetyl-4-methyl-5-oxo-hexanoic acid hexyl ester,

4-Acetyl-4-methyl-5-oxo-hexanoic acid octyl ester,
 4-Acetyl-4-methyl-5-oxo-hexanoic acid cyclopentadienyl ester,
 4-Acetyl-4-methyl-5-oxo-hexanoic acid (2-hydroxyethyl) ester,
 4-Acetyl-4-methyl-5-oxo-hexanoic acid cyclopentadienyl ester,
 5 4-Acetyl-4-methyl-5-oxo-hexanoic acid isobornyl ester,
 5-Acetyl-5-methyl-6-oxo-heptanoic acid ethyl ester,
 Diethyl 2,2-diacetyl-1,5-pentanedioate represented by the
 following formula,



10 3,3-Diacetyl-hexanedioic acid diethyl ester represented by the
 following formula,



4,4-Diacetyl heptanedioic acid 1,7-diethyl ester

15 4,4-Di-(1-oxopropyl)-heptanedioic acid 1,7-dimethyl ester

3,3-Diacetyl-1,5-dicyanopentane,

5,5-Diacetyl nonane-2,8-dione,

4,4-Diacetyl-heptanedioic acid 1,7-di-tert-butyl ester

4,4-Diacetyl-1,7-heptanedioic acid

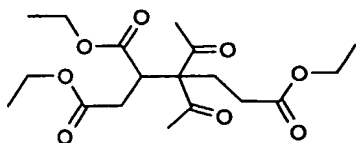
20 3,3-Diacetyl-1,5-bis(methylsulfonyl)-pentane

4,4-Diacetyl-7-oxo-octanoic-ethyl ester

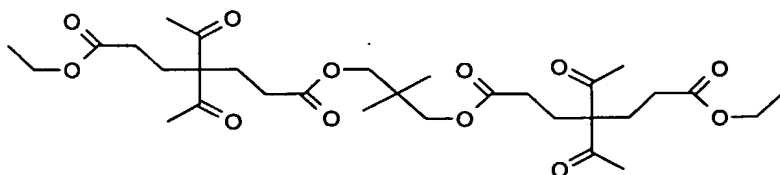
4,4-Diacetyl-5-(ethoxycarbonyl)-heptanedioic acid diethyl

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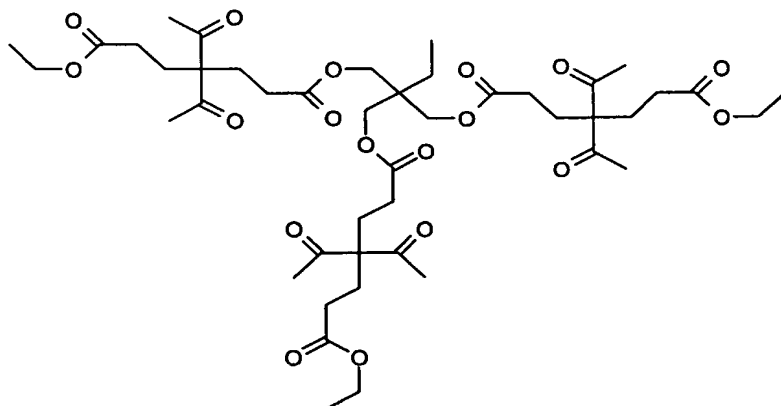
ester represented by the following formula,



a compound having two β -dicarbonyl groups such as 3-{4-Acetyl-
 5 4-[2-(ethoxycarbonyl)ethyl]-5-oxohexanoyloxy}-2,2-
 dimethylpropyl ethyl 4,4-diacetylheptane-1,7-dioate
 represented by the following formula,



10 a compound having three β -dicarbonyl groups such as 2,2-
 Bis({4-acetyl-4-[2-(ethoxycarbonyl)ethyl]-5-
 oxohexanoyloxy)methyl)butyl ethyl 4,4-diacetylheptane-1,7-
 dioate represented by the following formula,



15

Examples of the compound having the chemical structure

represented by formula (3) are

2-Acetyl-2-ethyl-pentanedioic acid 1,5-diethyl ester,

Diethyl 2-acetyl-2-(ethoxycarbonyl)-1,5-pentanedioate

3-Acetyl-3-(ethoxycarbonyl)-pentanedioic acid 1,5-dimethyl

5 ester

2-Acetyl-2-(ethoxycarbonyl)-hexanedioic acid 1,6-dimethyl

ester

4-Acetyl-4-(methoxycarbonyl)-heptanedioic acid 1,7-dimethyl

ester,

10 4-(Methoxycarbonyl)-4-(1-oxopropyl)-heptanedioic acid 1,7-

dimethyl ester,

4-(Ethoxycarbonyl)-4-(2-methyl-1-oxopropyl)-heptanedioic acid-

1,7-dimethyl ester

4-(Ethoxycarbonyl)-4-(1-oxobutyl)-heptanedioic acid 1,7-

15 dimethyl ester

4-Acetyl-4-(methoxycarbonyl)-octanedioic acid 1,8-dimethyl

ester

4-Acetyl-4-(tert-butoxycarbonyl)-heptanedioic acid 1,7-

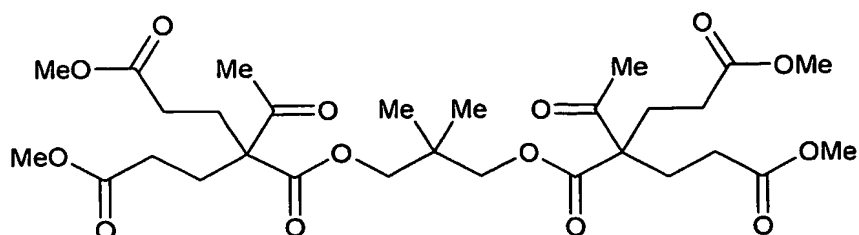
dimethyl ester

20 a compound having two β -dicarbonyl groups such as Dimethyl 4-

acetyl-4-[(3-{2,2-bis[2-(methoxycarbonyl)ethyl]-3-

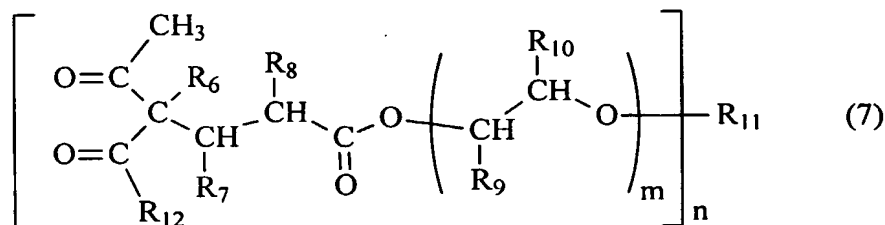
oxobutanoyloxy}-2,2-dimethylpropyl)oxycarbonyl]heptane-1,7-

dioate represented by the following formula,



Among these examples, the compounds having the chemical structure represented by formula (1) are preferred due to their excellent photosensitivity. Furthermore, the compounds having identical substituents at α -position of β -dicarbonyl structure are particularly preferred as well. From the viewpoint of ease of production, the compounds only having the above-mentioned γ position substituted group Nos. 4, 5, 12 are particularly preferred.

The novel compound of the present invention has the chemical structure represented by the following formula (7).



In these formulas, R_6 is selected from the group consisting of an alkyl group having a carbon number of 1 to 8, a C1-4 alkyl carbonyl group, a cyano group, a C1-4 alkyl carbonyl methyl group, a C1-4 alkyl carbonyl ethyl group, a C1-4 alkoxy carbonyl methyl group, a C1-4 alkoxy carbonyl ethyl group, and a alkyl group having a carbon number of 1 to 8 which is substituted by carboxyl group or cyano group.

The definition of the alkyl group having a carbon number of 1 to 8 is the same as the definition of the above-mentioned

R₃ or R₄.

The C1-4 alkyl carbonyl group includes acetyl group, ethyl carbonyl group, propyl carbonyl group, or butyl carbonyl group.

5 The C1-4 alkyl carbonyl methyl group includes 2-oxopropyl group, 2-oxobutyl group, 2-oxopentyl group, and 2-oxohexyl group.

The C1-4 alkyl carbonyl ethyl group includes 3-oxobutyl group, 3-oxopentyl group, 3-oxohexyl group, and 3-oxoheptyl
10 group. The C1-4 alkoxy carbonyl methyl group includes methoxy carbonyl methyl group, ethoxy carbonyl methyl group, propoxy carbonyl methyl group, butoxy carbonyl methyl group.

The C1-4 alkoxy carbonyl ethyl group includes methoxy carbonyl ethyl group, ethoxy carbonyl ethyl group, propoxy
15 carbonyl ethyl group, butoxy carbonyl ethyl group.

The alkyl group having a carbon number of 1 to 8 which is substituted by carboxyl group or cyano group includes carboxyl group- or cyano group-substituted methyl groups, ethyl groups, propyl groups, isopropyl groups, n-butyl groups, t-butyl
20 groups, pentyl groups, hexyl groups, cyclohexyl groups, n-heptyl groups, n-octyl groups.

R₇, R₈, R₉, and R₁₀ independently denote hydrogen atom, or a methyl group. Here, at least one of R₉ or R₁₀ is a halogen atom.

25 R₁₁ denotes a di-, tri-, tetra-valent aliphatic hydrocarbon group having a carbon number of 2 to 12. Examples of the R₁₁ are ethylene, propylene, trimethylol propane

residual group, and pentaerythritol residual group. Here, n is an integer of 2 to 4, and m is an integer of 0 to 15.

Furthermore, R_{12} denotes a methyl group, or an alkoxy group having a carbon number of 1 to 18.

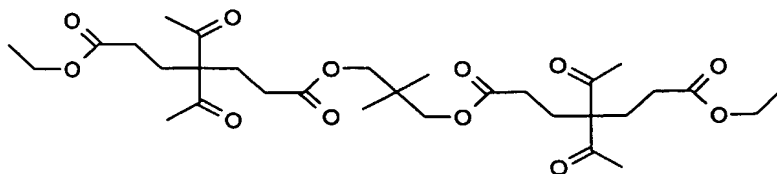
5 Examples of the alkoxy group include a linear or branched alkoxy group such as methoxy, ethoxy, propoxy, isopropoxy, n -butoxy, t -butoxy, pentyl-oxy, hexyl-oxy, cyclohexyl-oxy, n -heptyl-oxy, n -octyl-oxy, decyl-oxy, undecyl-oxy, lauryl-oxy, tridecyl-oxy, myristyl-oxy, pentadecyl-oxy, 10 cetyl-oxy, heptadecyl-oxy, and stearyl-oxy.

Among these novel compounds, the compounds having a molecular weight of 1000 or less are preferable as well as the above-mentioned photoinitiators from the viewpoints of compatibility and the cured product properties. Furthermore, 15 it is particularly preferred that the novel compounds have the chemical structure element represented by formula (2) at the rate of 17% to 54% by mass from the viewpoint of its photosensitivity if R_{12} is a methyl group. On the other hand, if R_{12} is an alkoxy group having a carbon number of 1 to 18, 20 the range of 17% to 47% by mass of the content of the chemical structure element represented by formula (2) is preferable.

Moreover, in the chemical structure represented by formula (7) or (8), formulas of $m=0$ are preferable from the viewpoints of balance of their molecular weight and the 25 content of the chemical structure element represented by formula (2).

Some of the above-mentioned examples belong to the

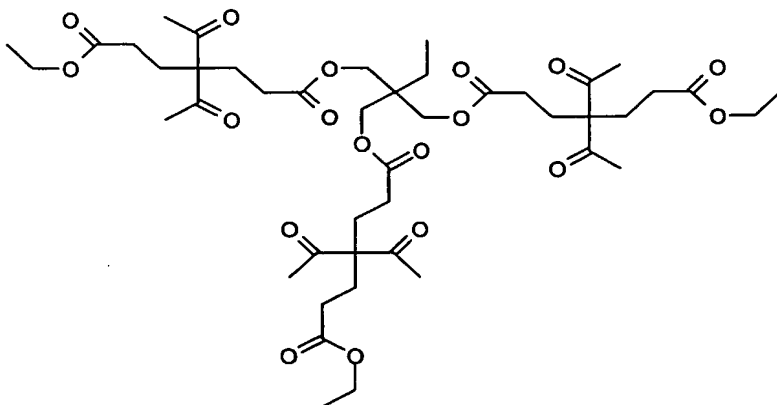
preferable novel compounds of the present invention such as 3-(4-Acetyl-4-[2-(ethoxycarbonyl)ethyl]-5-oxohexanoyloxy)-2,2-dimethylpropyl ethyl 4,4-diacetylheptane-1,7-dioate represented by the following formula,



5

a compound having three β -dicarbonyl groups such as 2,2-Bis(({4-acetyl-4-[2-(ethoxycarbonyl)ethyl]-5-oxohexanoyloxy)methyl})butyl ethyl 4,4-diacetylheptane-1,7-dioate represented by the following formula,

10



These compounds can be prepared by standard well-known organic syntheses, such as alkylation and Michael addition.

15 The alkylation can be carried out by the reaction of a β -diketone or a β -ketoester having an acidic H atom with a halogenated alkane in the presence of a base catalyst such as potassium carbonate. The Michael addition can be carried out

by the reaction of a β -diketone or a β -ketoester having an acidic H atom with an ethylenic unsaturated compound having an electron attracting group in the presence of a base catalyst.

Among them, the Michael addition is particularly suitable
5 for producing the compounds of the present invention.

Examples of suitable ethylenic unsaturated compounds having an electron attracting group, hereinafter abbreviated to "activated vinyl compound", as a starting material of the Michael addition are, for example, acrylic acid esters,
10 methacrylic acid esters, acryl amide, N-vinyl pyrrolidone or acrylonitrile.

Furthermore, specific examples thereof include, acrylic acid methyl ester, acrylic acid ethyl ester, acrylic acid butyl ester, acrylic acid hexyl ester, acrylic acid 2-ethyl
15 hexyl ester, acrylic acid octyl ester, acrylic acid ethyl ester cyclopentadienyl ester, acrylic acid isobornyl ester, acrylic acid 2-hydroxyethyl ester, acrylic acid 2-hydroxypropyl ester, propylenglykol mono acrylic acid ester, acrylic acid 2-carboxyethyl ester, acrylonitrile, vinyl methyl
20 ketone and acryl amide.

Furthermore, in the case of producing a compound having at least two β -diketone units per molecule, preferable examples of activated vinyl compound include

1,2-ethanediol diacrylate, 1,2-propanediol diacrylate,
25 1,4-butanediol diacrylate, hexan-1,6-diol diacrylate, dipropylene glycol diacrylate, neopentyl glycol diacrylate, ethoxylated neopentyl glycol diacrylate, propoxylated

neopentyl glycol diacrylate, tripropylene glycol diacrylate, polyethylene glycol diacrylate, trimethylolpropane triacrylate, ethoxylated trimethylolpropane triacrylate, propoxylated trimethylolpropane triacrylate, propoxylated glycerine triacrylate, tris (2-acryloyloxy ethyl) isocyanurate, pentaerythritol triacrylate, ethoxylated pentaerythritol triacrylate, pentaerythritol tetraacrylate, ethoxylated pentaerythritol tetraacrylate, di (trimethylolpropane) tetraacrylate, di (pentaerythritol) pentaacrylate, di (pentaerythritol) hexaacrylate.

Examples of the β -diketone or β -ketoester having an acidic H atom, typically have two carbonyl groups at the 1,3-position and one or preferably two acidic protons at the α position.

Examples of the compounds, which are suitable to synthesize the compounds related to this invention, are pentane-2,4-dione, hexane-2,4-dione, heptane-2,4-dione, 1-methoxy-pentan-2,4-dion, propionyl acetic acid ethyl ester, propionyl acetic acid butyl ester, butyryl acetic acid methyl ester, acetoacetic acid methyl ester, acetoacetic acid ethyl ester, acetoacetic acid isopropyl ester, acetoacetic acid butyl ester, acetoacetic acid tert-butylester, acetoacetic acid 2-methoxyethyl ester, acetoacetic acid 2-ethylhexyl ester, acetoacetic acid lauryl ester, 2-acetoacetoxyethyl acrylate, 2-acetoacetoxyethyl methacrylate and acetoacetic acid benzyl ester.

Furthermore, in the case of producing a compound having

at least two β -diketone units per molecule, preferable examples of activated vinyl compound include neopentyl glycol diacetoacetate, 2-ethyl-2-butyl-1,3-propanediol diacetoacetate, cyclohexanedimethanol acetoacetate, 5 trimethylolpropane triacetoacetate, glycerine triacetoacetate, pentaerythritol triacetoacetate, pentaerythritol tetraacetoacetate, ditrimethylolpropane tetraacetoacetate, and dipentaerythritol hexaacetoacetate.

Furthermore, the β -ketoester includes a β -ketoester of an 10 alkoxy alcohol obtained by a reaction of methanol, ethanol, propanol, isopropanol, n-butanol, t-butanol, pentanol, hexanol, cyclohexanol, n-heptanol, n-octanol, ethylene glycol, propylene glycol, trimethylol propane, or pentaerythritol and ethylene oxide.

15 The Michael addition reaction of the activated vinyl compound with the β -diketone or β -ketoester having an acidic H atom can be accelerated by special catalysts like strong bases and ammonium halides. The amount of catalyst used is within the range of 0.1 to 5.0 weight percent referred to the 20 complete reaction mixture, preferably 0.4 to 2.0 weight percent. Reaction temperature is within the range of 25 to 150°C, preferably 50 to 110°C. During the reaction of the activated vinyl compounds with the β -diketone or β -ketoester having an acidic H atom, the reaction mixture is sparged with 25 air.

Additionally, a polymerization inhibitor can be used in order to avoid unwanted gelation caused by reactive vinyl

compounds during synthesis. The amount is within the range of 0.01 to 0.5 weight percent. Suitable polymerization inhibitors are, for example, 4-methoxy phenol, phenothiazin and hydrochinone. By the above described procedure, obtained
5 products are either solid or liquid and can be purified by re-crystallization or distillation under reduced pressure.

The final product of the above-mentioned process is mainly obtained as a compound having a single, well-defined chemical structure. However, when using a poly-functional
10 activated vinyl compound or a poly-functional β -diketone or β -ketoester mentioned above, the final product may be obtained as a mixture containing several structure compounds.

The photoinitiator according to this invention is usable for curing of radical curable monomers, oligomers, or
15 polymers.

That is, the photocurable composition according to the present invention comprises,

- (i) an above-mentioned photoinitiator, and
- (ii) a radical curable ethylenic unsaturated compound

20 An amount of the photoinitiator among the photocurable composition is preferably within the range of 1 to 15 percent by weight. In particular, the amount is preferably within the range of 2 to 10 percent by weight.

Among the radical curable ethylenic unsaturated
25 compounds, especially those which are activated by heteroatoms can be polymerized very well. Examples for this type of compounds are monomers such as acrylic acid esters,

methacrylic acid esters, acrylonitrile, acryl amide, acrolein, vinyl acetate, vinyl propionate, N-vinyl pyrrolidone, N-vinyl carbazole, vinyl chloride and vinylidene chloride.

When using these monomers, the present composition can be
5 obtained by dissolving the photoinitiator in the monomers.
This is done by stirring or heating the mixture.

Furthermore, the radical curable ethylenic unsaturated compound also includes a compound having two or more radical curable ethylenic unsaturated groups. Examples for these
10 compounds are 1,2-ethanediol diacrylate, 1,2-propanediol diacrylate, 1,4-butanediol diacrylate, hexan-1,6-diol diacrylate, dipropylene glycol diacrylate, neopentyl glycol diacrylate, ethoxylated neopentyl glycol diacrylate, propoxylated neopentyl glycol diacrylate, tripropylene glycol
15 diacrylate, bisphenol A diglycidylether diacrylate, ethoxylated bisphenol A diglycidylether diacrylate, polyethylene glycol diacrylate, trimethylolpropane triacrylate, ethoxylated trimethylolpropane triacrylate, propoxylated trimethylolpropane triacrylate, propoxylated
20 glycerine triacrylate, tris(2-acryloyloxy ethyl) isocyanurate, pentaerythritol triacrylate, ethoxylated pentaerythritol triacrylate, pentaerythritol tetraacrylate, ethoxylated pentaerythritol tetraacrylate, di(trimethylolpropane) tetraacrylate, di(pentaerythritol) pentaacrylate,
25 di(pentaerythritol) hexaacrylate and oligomers and polymers containing acrylate groups obtained by conversion of poly epoxides with acrylic acid (epoxy acrylate) or by conversion

of polyester polyol with acrylic acid or monomeric alkyl acrylates (polyester acrylates) or by conversion of isocyanate prepolymers with 2-hydroxyethyl acrylate ((polyurea acrylate) and acrylated soy bean oil and acrylated silicone oil.

5 The composition of the present invention may contain another conventional photoinitiator as long as the effects of the present invention are not impaired.

 The conventional photoinitiator includes benzophenone, Michler's ketone, dialkyl acetophenone, hydroxyalkyl
10 acetophenone, aminoalkyl phenone, acylphosphine oxide and so called sensitizers such as isopropyl thioxanthone and 3-keto cumarine.

 Additionally the photocurable composition of the invention may contain so called accelerators such as
15 tributylamine, N-methyl diethanolamine, N-butyl diethanolamine, triethanolamine, piperidine, morpholine, piperazine, and acrylated amines, obtained from 1,6-hexanediol diacrylate and ethanolamine.

 In order to prevent inhibition of polymerization by
20 oxygen, a waxy compound can be added in addition to the above components. In consequence of their appropriate solubility in the mixtures, they float on top of the mixture at the start of polymerization and form a thin protecting layer between atmospheric oxygen and the polymerizing mixture.

25 Additionally, auto-oxidizing compounds like allyl ethers can be added that prevent inhibition of polymerization by oxygen in some cases.

Furthermore, the photocurable composition may contain well known additives, for example, silica, cement, talcum, leveling agents, wetting agents like polyelectrolyte, degassers like poly siloxane copolymers, flow and fluxing agents, surfactants, delustering agents, and plasticizers such as a phthalate.

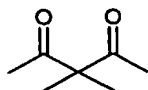
The photocurable composition according to this invention can be obtained by mixing the foregoing respective components uniformly, and can be cured by high-energy rays, preferably ultraviolet rays.

As radiation sources, sunlight or artificial irradiation generated by commercial mercury high-, medium- or low-pressure lamps, or xenon or wolfram lamps, can be used. The wavelength of the applied irradiation is within a range of 200 to 500 nm, preferably 250 to 350 nm. Duration of exposure depends on amount and type of the used photoinitiator and can be selected from a fraction of a second to several minutes. In mass polymerization, exposure time can be within the range of hours as well.

Examples

Example 1:

3,3-Dimethyl-2,4-pentandione



50 g of 2,4-pentandione, 178 g of iodo-methane, 120 g of

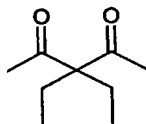
potassium carbonate, and 340 ml of acetone were mixed and heated to reflux for 18 hours. After cooling, 200 ml of petroleum ether was added, and the solid was filtered off and washed with 300 ml of an one-to-one mixture of petroleum ether. Residual 3-methyl-2,4-pentandione was removed by reacting with ethyl acrylate. The raw product was distilled to yield 23 g of 3,3-dimethyl-2,4-pentandione (boiling point: 63°C/20 mbar; purity: > 99%; colorless).

Content of the active structural element C₃O₂: 54%.

10

Example 2

3,3-Diethyl-2,4-pentandione



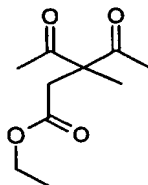
25 g of 2,4-pentandione, 98.3 g of bromo-ethane, 80 g of potassium carbonate, and 175 ml of acetone were mixed and refluxed for 39.5 hours. The solid was filtered off and washed with petroleum ether and acetone. To the liquid layer, 2.0 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene was added and the mixture was refluxed for two days. After addition of an additional 11.6 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene and refluxing for seven hours, 25 g of ethyl acrylate was added and the mixture was stirred at room temperature, and the solvent was distilled off and the mixture was refluxed for 8 hours. The base was neutralized by adding acetic acid, washed with an aqueous potassium carbonate solution, and extracted

with diethyl ether. The ether layer was dried and the product was isolated by fractional distillation to yield 9 g of product (boiling point 190°C/1000 mbar; purity: 75%; colorless).

5 Content of the active structural element C₃O₂: 44%.

Example 3

3-Acetyl-3-methyl-4-oxo-pentanoic acid ethyl ester



10 28 g of potassium hydroxide dissolved in 28 ml of water was added to a mixture of 50 g of 2,4-pentandione, and 100 ml and 115 ml of dioxane at 0 to 10°C. After 20 minutes of stirring, 125 g of bromoacetic acid ethyl ester was added dropwise and the mixture was stirred for 19 hours. The layers
15 were separated and washed with water and diethyl ether. The organic layer was dried and 3-acetyl-4-oxo-pentanoic acid ethyl ester was isolated by fractional distillation.

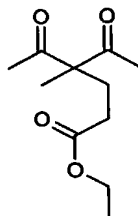
 48.3 g of 3-acetyl-4-oxo-pentanoic acid ethyl ester, 41 g of iodo-methane, 34 g of potassium carbonate, and 90 ml of
20 acetone were mixed and refluxed for 9 hours. After cooling, 100 ml of petroleum ether was added, and the solid was filtered off and washed with acetone and petroleum ether. Fractional distillation yielded 33.4 g of 3-acetyl-3-methyl-4-oxo-pentanoic acid ethyl ester (boiling point:

69°C/0.02 mbar; purity: 97%; colorless).

Content of the active structural element C₃O₂: 34%.

Example 4

5 4-Acetyl-4-methyl-5-oxo-hexanoic acid ethyl ester



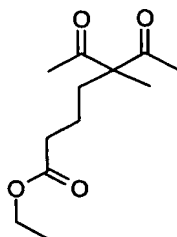
51 g of 2,4-pentandione, 89 g of iodo-methane, 66 g of potassium carbonate, and 170 ml of acetone were refluxed for 7 hours, the mixture was cooled, and 200 ml of petroleum ether
10 was added. After filtration, the solid was washed with acetone and petroleum ether, and 3-methyl-2,4-pentandione was isolated by fractional distillation.

30 g of 3-methyl-2,4-pentandione and 0.4 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were mixed and 39.3 g of
15 ethyl acrylate was added dropwise with the temperature being controlled by cooling with ice. Stirring was continued for 16 hours, the base was neutralized, 50 ml of diethyl ether was added, and the mixture was extracted with an aqueous sodium carbonate solution and water. Fractional distillation of the
20 organic layer yielded 33.4 g of 4-acetyl-4-methyl-5-oxo-hexanoic acid ethyl ester (boiling point: 95°C/0.02 mbar; purity: 98%; colorless).

Content of the active structural element C₃O₂: 32%.

Example 5

5-Acetyl-5-methyl-6-oxo-heptanoic acid ethyl ester



5 10 g of sodium acetylacetonate and 20 g of potassium carbonate in 50 ml of acetone were heated to reflux and 15 g of bromoacetic acid ethyl ester was added dropwise over 15 minutes. After refluxing for several days, the solid was filtered off and washed with petroleum ether and acetone.

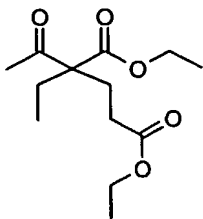
10 Fractional distillation yielded 6.4 g of 5-acetyl-6-oxo-heptanoic acid ethyl ester.

5.5 g of 5-acetyl-6-oxo-heptanoic acid ethyl ester, 5.7 g of potassium carbonate, 15 g of iodo-methane, and 15 ml of acetone were refluxed for 18 hours. After cooling, petroleum ether was added, and the solid was filtered off and washed with acetone and petroleum ether. Fractional distillation and re-distillation yielded 5.11 g of 5-acetyl-5-methyl-6-oxo-heptanoic acid ethyl ester (boiling point: 151°C/20 mbar; purity: 82%; colorless).

20 Content of the active structural element C₃O₂: 30%.

Example 6

2-Acetyl-2-ethyl-pentanedioic acid 1,5-diethyl ester



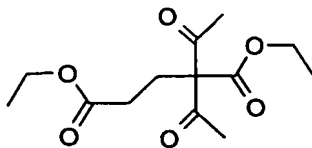
10 g of 2-ethyl ethyl acetoacetate and 0.1 g of 1,8-
5 diazabicyclo-[5.4.0]-undec-7-ene were mixed and 8 g of ethyl
acrylate was added at room temperature over 25 minutes. The
mixture was stirred for 23 hours, neutralized, poured into 20
ml of diethyl ether, and extracted with an aqueous potassium
carbonate solution and water. Fractional distillation of the
10 organic layer yielded 7.5 g of 2-acetyl-2-ethyl-pentanedioic
acid diethyl ester (boiling point: 145°C/20 mbar; purity:
>99%; colorless).

Content of the active structural element C₃O₂: 27%.

15

Example 7

Diethyl 2,2-diacetyl-1,5-pentanedioate



5 g of sodium salt of ethyl acetoacetate was mixed with
30 ml of toluene. Over 20 minutes, 5 g of acetic acid
20 chloride was added dropwise, and the mixture was stirred for
one hour at room temperature and at 85°C for 3 hours. The
solid was filtered off and washed with diethyl ether.

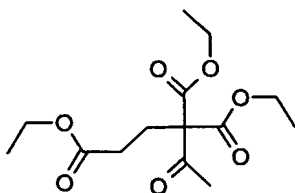
Fractional distillation of the liquid phase yielded ethyl 2-acetyl-3-oxo-butyrate.

2.0 g of ethyl 2-acetyl-3-oxo-butyrate was mixed with 5 drops of 1,8-diazabicyclo-[5.4.0]-undec-7-ene. Over 10 minutes, 2.7 g of ethyl acrylate was added dropwise. The mixture was stirred for 30 hours at room temperature, 2 ml of acetic acid was added to neutralize the base, and the mixture was extracted with an aqueous sodium carbonate solution and water. Fractional distillation yields 1.5 g of 2,2-diacetyl-1,5-pentanedioic acid diethyl ester (purity: 70%; colorless).

Content of the active structural element C_3O_2 : 25%.

Example 8

Diethyl 2-acetyl-2-(ethoxycarbonyl)-1,5-pentanedioate



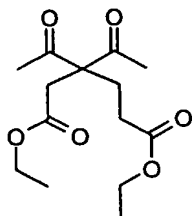
15

60 g of diethyl 2-carboxyethyl-1,5-pentanoate was added drop by drop to 10.44 g of sodium hydride covered by 100 ml of toluene. When heat-production ceased, 28 g of ethyl chloroformate was added dropwise. When maximum conversion was reached, the pH value was adjusted to 7 by adding hydrochloric acid, the salt was filtered off and the residual liquid fractionated under reduced pressure to yield diethyl 2-acetyl-2-(ethoxycarbonyl)-1,5-pentanedioate (purity: 85%; colorless).

Content of the active structural element C₃O₂: 23%.

Example 9

3,3-Diacetyl-hexanedioic acid diethyl ester



5

37 g of 3-acetyl-4-oxo-pentanoic acid ethyl ester, synthesized according to the procedure given in Example 15, and 0.3 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were mixed, and 30 g of ethyl acrylate was added dropwise over 30 minutes.

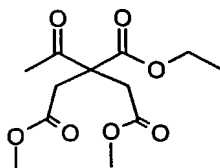
10 After stirring for 24 hours, the base was neutralized by adding acetic acid. 80 ml of diethyl ether was added, and the mixture was extracted with 60 ml of an aqueous sodium carbonate solution and 150 ml of water. Fractional distillation of the water-free organic layer yielded 46.2 g of
15 3,3-diacetyl-hexanedioic acid diethyl ester (boiling point: 150°C/0.03 mbar; purity: > 99%; colorless).

Content of the active structural element C₃O₂: 24%.

32

Example 10

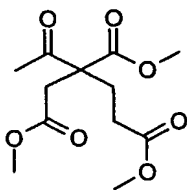
3-Acetyl-3-(ethoxycarbonyl)-pentanedioic acid 1,5-dimethyl ester



- 5 86 g of potassium *tert*-butylate was dissolved in 150 ml of dimethyl acetamide and 100 g of acetoacetic acid ethyl ester was added drop by drop with stirring. Subsequently, 83 g of chloroacetic acid methyl ester was added dropwise. When the exothermic reaction ended, 5 g of potassium *tert*-butylate
- 10 and 5 g of chloroacetic acid methyl ester were added alternately until conversion was complete. The solid was filtered off and washed with ethyl acetate. Fractional distillation yielded 30 g of 3-acetyl-3-(ethoxycarbonyl)-pentanedioic acid dimethyl ester (purity: 91%; colorless).
- 15 Content of the active structural element C₃O₂: 25%.

Example 11

2-Acetyl-2-(ethoxycarbonyl)-hexanedioic acid 1,6-dimethyl ester



20

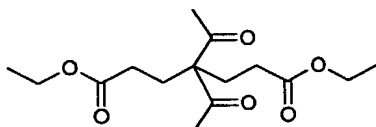
15 g of methyl acrylate was added dropwise to a mixture consisting of 30 g of 2-acetyl-1,4-butyric acid dimethyl ester

and 0.5 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene. The mixture was stirred at 75°C for 4 hours. After cooling, it was neutralized with acetic acid, washed with an aqueous potassium carbonate solution and was subsequently washed with water, and distilled, and colorless 2-acetyl-2-(ethoxycarbonyl)-hexanedioic acid dimethyl ester was obtained at 95% purity.

Content of the active structural element C₃O₂: 25%

10

Example 12



4,4-diacetyl heptanedioic acid 1,7-diethyl ester

20 ml of 2,4-pentandione, 40 ml of ethyl acrylate, and 1.5 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were mixed, and stirred for 3 hours at 80°C in air. The crude product was distilled (boiling point: 155°C/10⁻² mbar). The distilled product was crystallized in rhombic crystals having purity of 99.4% (gas chromatography). Melting point: 64°C. The product was colorless. Content of the active structural element C₃O₂: 23%.

20

Example 13

4-Acetyl-4-(methoxycarbonyl)-heptanedioic acid 1,7-dimethyl ester

50.0 g of methyl acetoacetate and 100.0 g of methyl

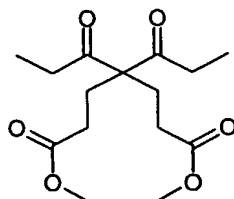
acrylate were mixed, and 0.6 g of tetramethyl guanidine and 0.05 g of 4-methoxy phenol were added. The mixture was stirred at 85°C for 3 hours in air, and the excess methyl acrylate was removed by distillation, leading to a yellowish liquid crude product with a purity of 91%. The final
5 fractional distillation of the crude product yielded the subject compound, which was colorless, having a boiling point: 145°C (10⁻² mbar) and purity: GC > 99%.

Content of the active structural element C₃O₂: 24%.

10

Example 14

4,4-Di-(1-oxopropyl)-heptanedioic acid 1,7-dimethyl ester

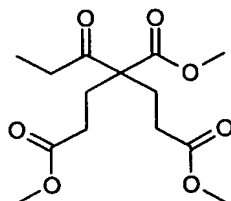


40.0 g (0.465 mol) of methyl acrylate was added drop by
15 drop to 21.9 g (0.171 mol) of 3,5-heptanedione and 0.2 g (1.3 × 10⁻³ mol) of DBU. After addition of two thirds of the acrylate, temperature was raised to 40°C. Stirring was continued for three hours, then the mixtures was heated to reflux, and an additional 0.5 g (3.3 × 10⁻³ mol) of DBU was
20 added. Refluxing was continued for 6 hours, excess acrylate was distilled off, and the product was isolated by fractional distillation. 22.8 g (0.08 mol) of dimethyl 4,4-di-(1-oxopropyl)-heptanedioate was obtained (boiling temperature: 175°C at 0.02 mbar; yield: 44%; colorless).

Content of the active structural element C_3O_2 : 23%.

Example 15

4-(Methoxycarbonyl)-4-(1-oxopropyl)-heptanedioic acid 1,7-
dimethyl ester

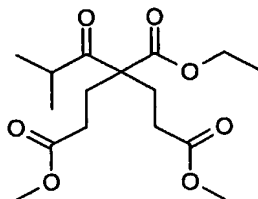


39.9 g (0.31 mol) of methyl 3-oxo-pentanoate and 1.9 g (0.012 mol) of DBU were mixed at room temperature in a three necked round bottom flask, equipped with a dropping funnel and a reflux condenser. 66.4 g (0.77 mol) of methyl acrylate was slowly added drop by drop. After the exothermal behavior has leveled off, the reaction mixture was stirred at 60°C for an additional two hours. Cyclohexane was added and the mixture was washed with diluted hydrochloric acid and was subsequently washed with distilled water until the aqueous layer showed a neutral reaction. The organic layer was dried over sodium sulfate, the solvent was removed, and the addition product was isolated by fractional distillation in a vacuum. 21.4 g (0.20 mol) of 4-(methoxycarbonyl)-4-(1-oxopropyl)-heptanedioic acid 1,7-dimethyl ester was obtained (yield: 64%; boiling temperature: 130°C at 0.02 mbar; colorless).

Content of the active structural element C_3O_2 : 23%.

Example 16

4-(Ethoxycarbonyl)-4-(2-methyl-1-oxopropyl)-heptanedioic acid-
1,7-dimethyl ester

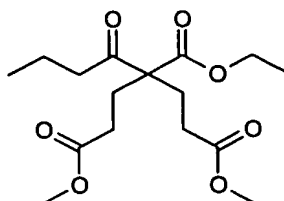


5 12.0 g (0.08 mol) of ethyl 4-methyl-3-oxo-pentanoate and
0.5 g (0.003 mol) of DBU were placed in a three necked round
bottom flask, equipped with dropping funnel and reflux
condenser. At room temperature, 16.0 g (0.18 mol) of methyl
acrylate was slowly added drop by drop. After the exothermal
10 behavior of the reaction leveled off, the reaction mixture was
stirred at 60°C for an additional two hours. The mixture was
poured into cyclohexane and was washed with diluted
hydrochloric acid and was subsequently washed with distilled
water until the aqueous layer was neutral. The organic layer
15 was dried over sodium sulfate, the solvent was removed, and
the addition product was isolated by fractional distillation
in a vacuum. 15.0 g (0.045 mol) of 4-(ethoxycarbonyl)-4-(2-
methyl-1-oxopropyl)-heptanedioic acid 1,7-dimethyl ester was
obtained (yield: 56%; boiling point: 145°C at 0.02 mbar;
20 colorless).

Content of the active structural element C₃O₂: 21%.

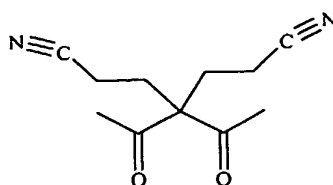
Example 17

4-(Ethoxycarbonyl)-4-(1-oxobutyl)-heptanedioic acid 1,7-dimethyl ester



5 41.2 g (0.26 mol) of ethyl 3-oxo-hexanoate and 1.9 g
(0.012 mol) of DBU were placed in a three necked round bottom
flask, equipped with dropping funnel and reflux condenser. At
room temperature, 56.3 g (0.65 mol) of methyl acrylate was
slowly added drop by drop. After the exothermal behavior of
10 the reaction leveled off, the reaction mixture was stirred at
60°C for an additional two hours. The mixture was poured into
cyclohexane and was washed with diluted hydrochloric acid and
was subsequently washed with distilled water until the aqueous
layer showed a neutral reaction. The organic layer was dried
15 with sodium sulfate, the solvent was removed, and the residual
addition product was isolated by fractional distillation in a
vacuum. 50.0 g (0.15 mol) of 4-(ethoxycarbonyl)-4-(1-
oxobutyl)-heptanedioic acid 1,7-dimethyl ester was obtained
(yield: 58%; boiling point: 149°C at 0.007 mbar; colorless).
20 Content of the active structural element C₃O₂: 21%.

Example 18



3,3-Diacetyl-1,5-dicyanopentane

After addition of 0.25 g of KOH, dissolved in 2.5 ml of methanol, to 15.0 g of 2,4-pentandione, 35.0 g of acrylonitrile was added dropwise with the reaction temperature not exceeding 40°C. The product which was precipitated during the process was filtered off, washed with a small amount of acetone, and dried. Melting point: 185-186°C. Purity > 98%. The product was colorless.

Content of the active structural element C₃O₂: 33%.

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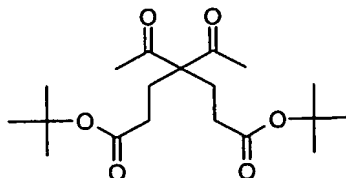
Example 19

5,5-Diacetyl nonane-2,8-dione

A mixture of 15.0 g of 2,4-pentandione and 0.35 g of tetramethyl guanidine was added to 40.0 g of 2-butanone drop by drop, with the reaction temperature not exceeding 40°C. Then, the excess of 2-butanone was removed by distillation under reduced pressure (200 mbar). The final product was obtained at 96% purity as a colorless product. Melting point: 59°C. Content of the active structural element C₃O₂: 28%.

20

Example 20

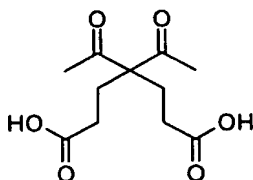
4,4-Diacetyl-heptanedioic acid 1,7-di-*tert*-butyl ester

At room temperature, 282 g of *tert*-butyl acrylate was
5 added dropwise to a mixture of 100 g of 2,4-pentandione and 3
g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene and stirred for 5
hours. A small portion of methanol was added to the mixture
at 40°C. A crude product was precipitated from the mixture at
room temperature and was recrystallized from methanol to
10 obtain 4,4-diacetyl-heptanedioic acid di-*tert*-butyl ester as
white crystals at a yield of 54% (purity: 100%; colorless).

Content of the active structural element C₃O₂: 20%.

Example 21

15 4,4-Diacetyl-1,7-heptanedioic acid



4,4-Diacetyl-heptanedioic acid di-*tert*-butyl ester was
hydrolyzed at 95°C over 5 hours using a mixture of *tert*-
butanol and water as solvent and 11 weight-% hydrochloric
20 acid. The solvent was removed and the obtained crude product
was recrystallized from a mixture of acetone and petroleum
ether (1:1) to yield 4,4-diacetyl-heptanedioic acid as white

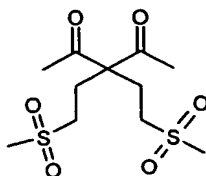
crystals at a yield of 69%. (melting point: 176°C; purity 94%; colorless).

Content of the active structural element C₃O₂: 28%.

5

Example 22

3,3-Diacetyl-1,5-bis(methylsulfonyl)-pentane



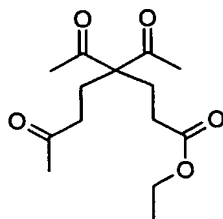
6 g of 2,4-pentandione and 3 drops of triethyl amine were mixed and 7 g of methyl vinyl sulfone was added dropwise at room temperature. After stirring at 60°C for 7 hours, 5 drops of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were added and the mixture was stirred at room temperature for 12 hours. The mixture was stirred with 25 ml of ethanol and neutralized after a white solid was precipitated. The mixture was poured into 200 ml of water and ice, and was kept in the refrigerator overnight. The solvent was filtered off, dried and recrystallized from 500 ml of ethanol. 2.9 g of white, long needles of the subject compound was obtained (melting point: 162°C; purity: >99%; colorless).

20

Content of the active structural element C₃O₂: 22%.

Example 23

4,4-Diacetyl-7-oxo-octanoic-ethyl ester



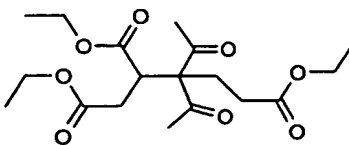
60 g of 2,4-pentandione and 0.1 g of triethyl amine were
5 mixed, and 28 g of 2-butanone was added dropwise at room
temperature. The mixture was stirred for 42 hours and the raw
product was distilled off. Re-distillation gave 31.4 g of 3-
acetyl-2,6-heptandione.

10.2 g of 3-acetyl-2,6-heptandione and 5 drops of 1,8-
10 diazabicyclo-[5.4.0]-undec-7-ene were mixed, and 9 g of ethyl
acrylate was added dropwise over 15 minutes. After stirring
for 16 hours, the base was neutralized with acetic acid, 30 ml
of diethyl ether was added, and the solution was extracted
with sodium carbonate solution and water. Fractional
15 distillation of the dry organic layer yielded 9.3 g of 4,4-
diacetyl-7-oxo-octanoic-ethyl ester (boiling point:
136°C/0.02 mbar; purity: 81%; colorless).

Content of the active structural element C₃O₂: 26%.

Example 24

4,4-Diacetyl-5-(ethoxycarbonyl)-heptanedioic acid diethyl ester

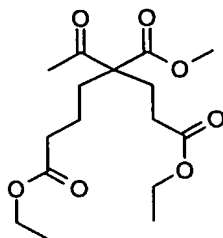


5 a) To a mixture of 70 g of 2,4-pentandione and 1.72 g of trioctyl phosphine, 45 g of ethyl acrylate was added dropwise over 40 minutes, keeping the temperature at 0-5°C by cooling. After stirring for 65 hours, 2 ml of acetic acid and 150 ml of diethyl ether were added, and the solution was extracted with
10 an aqueous sodium carbonate solution and water. The organic layer was dried, and 4-acetyl-5-oxohexanoic acid ethyl ester was isolated by fractional distillation.

b) 25.7 g of 4-acetyl-5-oxohexanoic acid ethyl ester and 0.3 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were mixed, and
15 35 g of fumaric acid diethyl ester was added at room temperature over 40 minutes. After stirring for 24 hours, the amount of 1,8-diazabicyclo-[5.4.0]-undec-7-ene was doubled and stirring continued until conversion was complete. 0.5 g of acetic acid and 50 ml of diethyl ether were added, and the
20 solution was extracted with an aqueous potassium carbonate solution and water. Fractional distillation of the dry organic layer yielded 25.6 g of 4,4-diacetyl-5-(ethoxycarbonyl)-heptanedioic acid diethyl ester (boiling point: 170°C/0.002 mbar; purity: 93%; colorless). Content of
25 the active structural element C₃O₂: 18%.

Example 25

4-Acetyl-4-(methoxycarbonyl)-octanedioic acid 1,8-dimethyl ester



5

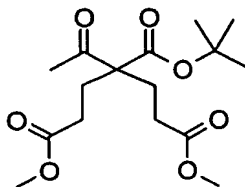
0.3 g of sodium was dissolved in 160 g of methyl acetoacetate, and 86 g of methyl acrylate was added drop by drop at a temperature below 30°C. The reaction was continued for one hour, and subsequently the pH was adjusted to 7. The salt was filtered off and the raw product purified by distillation to yield 2-acetyl pentanoic 5-ethyl-1-methyl ester.

12.5 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene and 2.3 sodium were dissolved in 50 g of 2-acetyl pentanoic 5-ethyl-1-methyl ester, and 48 g of bromobutyric acid ethyl ester was added drop by drop. When the temperature decreased, an additional 2 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene was added. After almost all educt was consumed, the salt was filtered off and 4-acetyl-4-(methoxycarbonyl)-octanedioic acid dimethyl ester was isolated by fractional distillation to yield 44 g of colorless product (purity: 94%).

Content of the active structural element C₃O₂: 21%.

Example 26

4-Acetyl-4-(tert-butoxycarbonyl)-heptanedioic acid 1,7-dimethyl ester

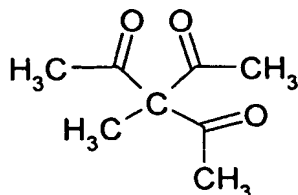


5 0.5 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene and 40 g of
tert-butyl acetoacetate were stirred at room temperature. 60
g of methyl acrylate was dropped in. The exothermic reaction
started immediately. The temperature of the mixture was
allowed to reach 80°C. The mixture was stirred at 80°C for an
10 additional two hours. The basic catalyst was removed by
washing with diluted hydrochloric acid and the organic layer
was fractionated in a vacuum. The pure colorless 4-acetyl-4-
(tert-butoxycarbonyl)-heptanedioic acid 1,7-dimethyl ester
boils at 149°C/0.025 mbar (yield: 56 g).

15 Content of the active structural element C₃O₂: 21%.

Example 27

1,1,1-Triacetylene



20 3-methylpentanedione-2,4 was prepared from acetylacetone
and methyl iodide in the presence of potassium carbonate

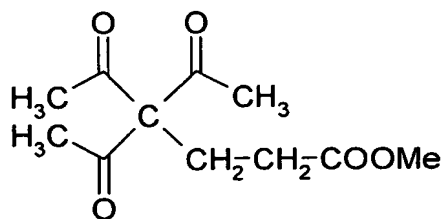
(Organic Syntheses, Coll. Vol. V, 785).

57 g of 3-methylpentanedione-2,4 was treated with 12 g of sodium hydride. During this process, the temperature was maintained at 30-40°C. Then, 200 ml of diethyl ether was added, and 39.25 g of acetyl chloride was dropped in. The mixture was stirred for 3 hours in boiling ether and was washed with potassium hydroxide. Diethyl ether was distilled off in a vacuum. The residual slightly yellowish liquid was the crude triacetyl ethane having a purity of above 90%.

Content of the active structural element C₃O₂: 44%.

Example 28

4,4-Diacetyl-5-oxohexanoic acid methyl ester



4-acetyl-5-oxohexanoic acid methyl ester was prepared through the Michael addition of methyl acrylate to acetyl acetone in the presence of metallic sodium. The crude product was purified by vacuum distillation and received in a purity of approximately 99%.

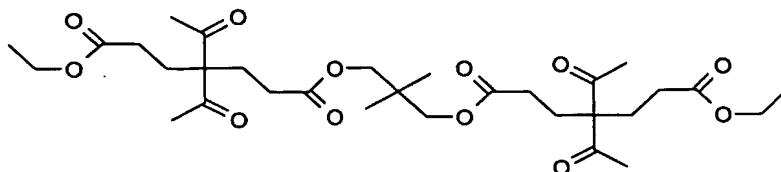
92.5 g of 4-acetyl-5-oxohexanoic acid methyl ester was reacted with 11.5 g of metallic sodium in 200 ml of diethyl ether. After the dissolving of sodium was finished, 39.25 g of acetyl chloride was dropped in, and the resulting boiling mixture was stirred for 6 hours. Then, the reaction mixture

was filtered, washed with aqueous potassium hydroxide and water, following by evaporation of solvent. The final vacuum distillation yielded the subject compound as a yellowish liquid having a purity of above 90%.

5 Content of the active structural element C_3O_2 : 30%.

Example 29

3-{4-Acetyl-4-[2-(ethoxycarbonyl)ethyl]-5-oxohexanoyloxy}-2,2-dimethylpropyl ethyl 4,4-diacetylheptane-1,7-dioate



10

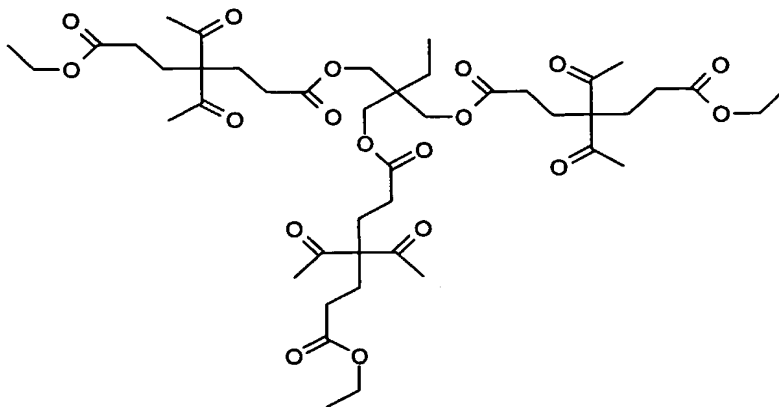
5.0 g of 4-acetyl-5-oxohexanoic acid ethyl ester and five drops 1,8-diazabicyclo-[5.4.0]-undec-7-ene were mixed and 2.6 g of neopentyl glycol diacrylate was added dropwise. The mixture was stirred for three hours to yield a highly viscous colorless oil. Gel permeation chromatography showed a main peak at 690 g/mol, nuclear magnetic resonance spectra and infrared spectrum are according to the expected structure.

15

Content of the active structural element C_3O_2 : 22%.

Example 30

2,2-Bis((4-acetyl-4-[2-(ethoxycarbonyl)ethyl]-5-oxohexanoyloxy)methyl)butyl ethyl 4,4-diacetylheptane-1,7-dioate



5

5.0 g of 4-acetyl-5-oxo-hexanoic acid ethyl ester and five drops 1,8-diazabicyclo-[5.4.0]-undec-7-ene were mixed, and 2.4 g of trimethylolpropane triacrylate was added dropwise. The mixture was stirred for three hours to yield a highly viscous colorless oil. Gel permeation chromatography showed a main peak at 958 g/mol, and nuclear magnetic resonance spectra and infrared spectrum were according to the expected structure.

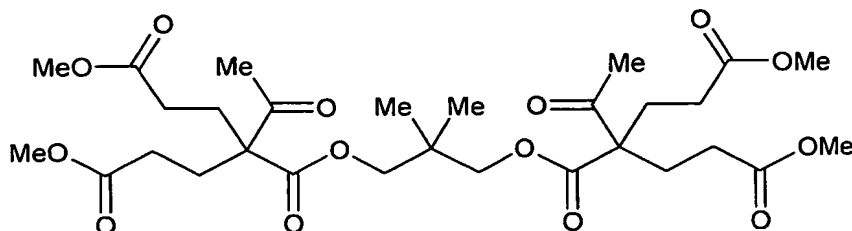
10

Content of the active structural element C_3O_2 : 23%.

15

Example 31

Dimethyl 4-acetyl-4-[(3-{2,2-bis[2-(methoxycarbonyl)ethyl]-3-oxobutanoyloxy}-2,2-dimethylpropyl)oxycarbonyl]heptane-1,7-dioate



5

Transesterification of neopentylglycol with ethyl acetoacetate yielded neopentylglycol diacetoacetate. The crude product was isolated through a short-path distillation (110°C at 0.001 mbar) to yield the pure (> 98%)

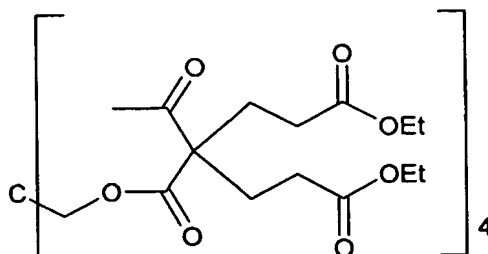
10 neopentylglycol diacetoacetate as a colorless liquid.

40 g of this diacetoacetate and 0.4 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were stirred at room temperature, and 100 g of methyl acrylate was added drop by drop. After the first exothermic reaction finished, the mixture was stirred for 3 hours at 80°C. Then, the excess of methyl acrylate was distilled off in a vacuum. The subject compound remained as a slightly yellowish viscous liquid. Yield was 88 g. Nuclear magnetic resonance spectra and infrared spectrum confirmed the expected structure.

20 Content of the active structural element C₃O₂: 22%.

Example 32

Photoinitiator based on pentaerythritol tetraacetoacetate and ethyl acrylate

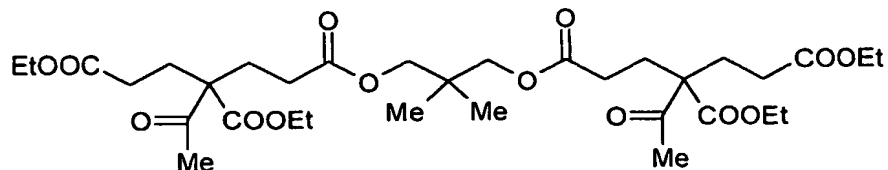


100 g of ethyl acrylate and 2 g of 10% methanolic KOH were added to 50 g of a tetra acetoacetate, synthesized by transesterification of pentaerythritol with ethyl acetoacetate. The mixture was stirred for 3 hours at 80°C. Subsequently the basic catalyst was neutralized with acetic acid. The resulting crude reaction product was a highly viscous, yellowish oil. Molecular weight $M_w = 860$ (GPC).

Content of the active structural element C_3O_2 : 21%.

Example 33

3-((4S)-4-(ethoxycarbonyl)-4-[2-(ethoxycarbonyl)ethyl]-5-oxohexanoyloxy)-2,2-dimethylpropyl ethyl (4S)-4-acetyl-4-(ethoxycarbonyl)heptane-1,7-dioate
(Photoinitiator based on NPGDA and acetyl diethylglutarate)



46 g of acetylglutaric acid diethyl ester and 0.5 g of

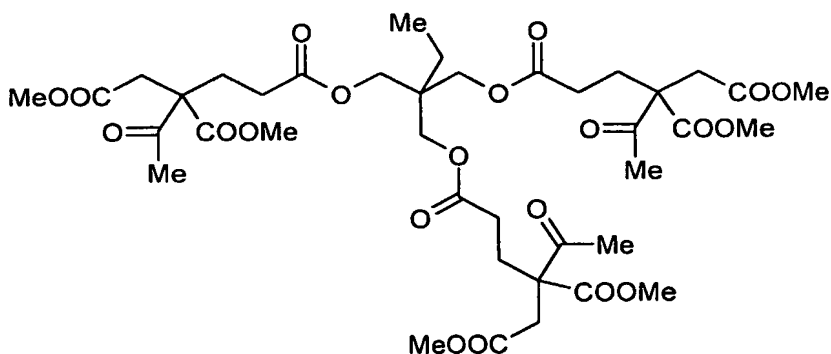
1,8-diazabicyclo-[5.4.0]-undec-7-ene were stirred at room temperature, and 21.2 g of neopentylglycol diacrylate was added drop by drop. After the first exothermic reaction finished, the mixture was stirred for 3 hours at 80°C. The subject compound remained as a slightly yellowish viscous liquid. Yield was 66.5 g. Nuclear magnetic resonance spectra and infrared spectrum confirmed the expected structure.

Content of the active structural element C₃O₂: 20%.

Example 34

2,2-Bis({4-(methoxycarbonyl)-4-[(methoxycarbonyl)methyl]-5-oxohexanoyloxy)methyl}butyl methyl 3-acetyl-3-(methoxycarbonyl)hexane-1,6-dioate

(Photoinitiator based on TMPTA and acetyl dimethylsuccinate)

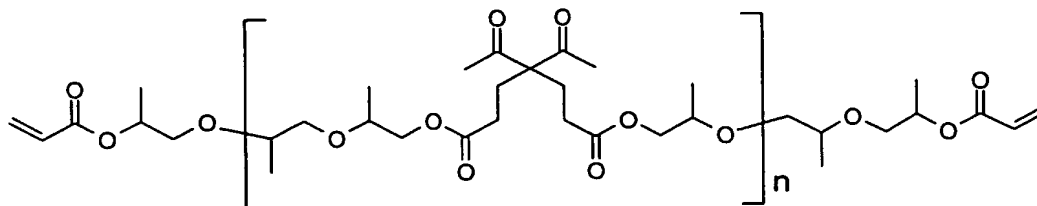


56.4 g of acetylsuccinic acid dimethyl ester and 0.5 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene were stirred at room temperature, and 30 g of trimethylolpropane triacrylate was added drop by drop. After the first exothermic reaction finished, the mixture was stirred for 3 hours at 80°C. The subject compound remained as a slightly yellowish viscous liquid. Yield was 86 g. Nuclear magnetic resonance spectra

and infrared spectrum confirmed the expected structure.

Content of the active structural element C_3O_2 : 24%.

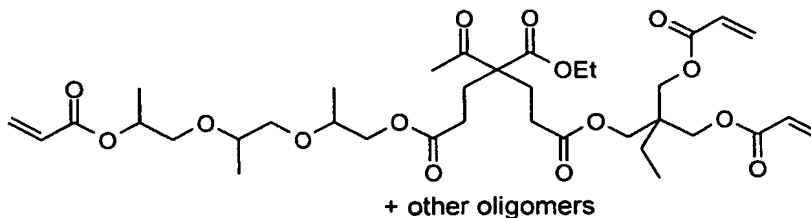
Comparative Example 1



0.10 mol of 2,4-pentandione, 0.195 mol of tripropylenglycol diacrylate, and 0.45 g of tetramethylguanidine were mixed and stirred under sparging with air. After the reaction had started, the temperature was allowed to reach 90°C. Then the mixture was cooled to 80°C and stirred at this temperature for one hour. After cooling, the product had a viscosity of 12000 mPas at 25°C and a molecular weight of $M_w = 2000$.

Content of the active structural element C_3O_2 : 9.86%.

Comparative Example 2



50.0 g of trimethylolpropane triacrylate, 50 g of tripropylene glycol diacrylate, and 15.0 g of ethyl acetoacetate were mixed, and this was heated to 50°C before 1.5 g of 1,8-diazabicyclo-[5.4.0]-undec-7-ene (DBU) was added.

The mixture was sparged with air. After the exothermic reaction started, the temperature was allowed to rise until 110°C. Then, the mixture was cooled to 80°C and stirred for 1 hour at 80°C. After cooling, the resulting product had a
5 viscosity of 18000 mPas at 25°C and a molecular weight of $M_w = 6000$.

Content of the active structural element C_3O_2 : 6.73%.

(Properties of Cured Coatings)

10 The following table shows properties of various cured coatings of photocurable compositions which contain photoinitiators from the above-mentioned examples.

Fist of all, photocurable compositions were prepared by mixing the composition in the table, then coated on aluminum
15 sheets as thin films and cured by UV irradiation.

The cured coatings were tested with respect to their pencil hardness and their solvent resistance towards ethyl methyl ketone.

Table:

System	Photoinitiator from example	concentration (weight%)	test substance	exposure energy ¹ 0.6 J/cm ²	solvent resistance ² > 75	pencil hardness ³ H-2H
1	12	3	UV-overprint varnish ⁴	0.6 J/cm ²	> 75	H-2H
2	12	7	UV-overprint varnish ⁴	0.6 J/cm ²	> 75	2H-3H
3	13	8	UV-overprint varnish ⁴	0.6 J/cm ²	~ 30	HB
4	12	5	epoxy acrylate ⁵	0.6 J/cm ²	> 75	3H
5	12	5	polyester acrylate ⁶	0.6 J/cm ²	> 75	H
6	13	10	polyester acrylate ⁶	0.6 J/cm ²	> 75	H
7 ⁸	18	3	epoxy acrylate ⁷	0.1J/cm ²	>75	4H
8	19	5	Acrylate mixture ⁹	0.5 J/cm ²	>100	2H

¹Quantity of light for crosslinking produced by a F 300H-lamp (Sum UV-A-B-C), measured with a radiometer from EIT.

²Solvent resistance of the cured film, tested by repeatedly rubbing the cured surface with a cotton sheet soaked in ethyl methyl ketone (MEK).

The number of recurrences, the so called double-rubs, performed without visible damage on the film was counted

³Hardness of pencil which caused visual damage on the cured film

⁴UV-overprint varnish, consisting of 40% tripropylene glycol diacrylate, 30% (trimethylolpropane) triacrylate and 30% bisphenol A diglycidylether diacrylate.

⁵Epoxy acrylate Ebecryl 120 from UCB.

⁶Polyester acrylate Ebecryl 835 from UCB.

⁷Epoxy acrylate, obtained by reacting Epiclone 1050 from DIC with acrylic acid.

⁸Curing was done at 110°C, as the curable mixture is solid at 25°C.

⁹Acrylate mixture comprising 30% by weight of Dipropylene glycol diacrylate (DPGDA), 45% by weight of an acrylate compound obtained by reacting Epiclone 1050 from DIC with acrylic acid, and 25% by weight of Trimethylolpropane triacrylate (TMPTA).

The photoactivity of all prepared photoinitiators was tested.

For UV-curing tests, a mixture consisting of 30 weight-% tripropylene glycol diacrylate (TPGDA), 25 weight-% trimethylolpropane triacrylate (TMPTA) and 45 weight-% Ebecryl 150 (UCB) was used. A solution of 5 weight-% of the tested photoinitiator in the UV-curable mixture was prepared and coated on an aluminum sheet as a film of 50 μm thickness. The sample was irradiated with a Fusion F 300 using the H-bulb (300 W/inch). The radiation doses were 0.087 J/cm² for UV-A, 0.058 J/cm² for UV-B and 0.035 J/cm² for UV-C, which gives 180 mJ/cm² in sum. The belt-speed was adjusted to 16 m/min and 3 passes were done.

All tested photoinitiators of the invention are photoactive and initiate a polymerization under irradiation, indicated by the formation of solvent-resistant films. The solvent resistance of the cured film was tested by rubbing the film with a cotton sheet soaked with 2-butanone (methyl ethyl ketone; MEK) and counting the double rubs needed to cause visible defects in the film. The results are summarized in the following table:

Photoinitiator according to example No.	Solvent resistance of the cured film (MEK double rubs)
1	> 100
2	> 100
3	> 100
4	~ 12
5	> 100
6	~ 13
7	~ 32
8	~ 28
9	> 100
10	> 100
11	> 100
14	~ 40
15	~ 42
16	~ 52
17	~ 20
19 ^{a)}	> 100
20	> 100
21 ^{b)}	> 100
22 ^{c)}	> 80
23	> 100
24	> 100
25	~ 40
26	~ 60
27	> 100
28	~ 14
29	> 100
30	> 100
31	~ 60
32	~ 70
33	< 80
34	< 100

- a) a test mixture consisting of 29% dipropylene glycol diacrylate (DPGDA), 42% Ebecryl 150 and 24% TMPTA was used
- b) a solution of 4 weight-% of the photoinitiator in a mixture of 20% deionized water and 76% Photomer 3165 (Cognis) was tested
- c) 10 weight-% of dimethyl sulfoxide were added to the test mixture in order to dissolve the photoinitiator

As a reference, a film of the same thickness was coated on the same sheet consisting of the UV-curable mixture, however, without a photoinitiator of the present invention. This film showed no curing and was easily removed by whipping with a dry cotton sheet.

Also, the influence of the molecular weight of the photoinitiators on their photoactivity was investigated.

Substrate: Aluminum

Film Thickness: 50 μm (wet)

¹Irradiance: 0.5 J/ cm^2

Bulb: Fusion F 300H bulb

Test-mixture: Standard UV over-print varnish:

Dipropyleneglycol diacrylate (DPGDA) 30%

Acrylated polyester Ebecryl 150 45%

Trimethylolpropane triacrylate (TMPTA) 25%

Procedure: The photoinitiators were dissolved in the test mixture. The test mixture was applied on an aluminum substrate at a film thickness of 50 μm (wet) and cured under UV-light with a Fusion F 300 H bulb. The applied irradiance was 0.5 J/ cm^2 . Then, the coatings were assessed by appearance, solvent resistance and hardness.

5

Entry	² PI-Type	% PI	³ Coating viscosity	Surface appearance	⁴ Solvent resistance	⁵ Pencil hardness	Molecular weight
1	-	-	200 mPas	wet	0	-	
2	A	5	200 mPas	dry	> 100	2H	240
3	B	5	200 mPas	dry	> 100	2H	300
4	C	5	260 mPas	wet	~ 5	< 6B	1800
5	C	13.7	290 mPas	wet-tacky	~ 25	HB	1800
6	C	24.2	340 mPas	tacky	> 100	H	1800
7	C	32.4	540 mPas	dry	> 100	H-2H	1800

¹Radiant energy of the UV-bulb (total UVA-C) at the surface of the coating measured with the radiometer UVICURE from EIT company.

²PI type:

10 A = 5,5-diacetylnonane-2,8-dione, (Example 19 of this invention)

B = diethyl-4,4-diacetylheptanedioate, (Example 12 of this invention)

15 C = photoactive resin prepared according to US 5945487, column 9, line 45.

³Coating viscosity, measured with an ICI cone and plate viscosimeter at a shear rate of 5000 D^{-1}

⁴Solvent resistance of the hardened film, tested by repeated rubbing of the film surface with a woodpulp cloth impregnated with methyl ethyl ketone (MEK). The number of rubbings that still did not produce any visible damage to the coating was measured.

(properties comparison with conventional photoinitiators)

Photoinitiator	Example 19	2-hydroxy-2-methylpropio phenone	Bisacylphenyl phosphineoxide
Color	Colorless	slightly yellow	slightly yellow
Molar extinction coefficients at 300 nm in methanol	122 ml/g·cm	822 ml/g·cm	18200 ml/g·cm

Due to the low molar extinction coefficient, photoinitiators of the present invention are particularly suitable for curing of thick coatings layers, which was demonstrated by the following experiment:

A container, having dimensions of 1 cm x 1 cm x 1 cm, was filled with a test mixture, containing 30% of trimethylolpropane triacrylate, 30% of bisphenol-A-diglycidylether diacrylate, 35% of dipropylene glycol diacrylate and 5% of 5,5-diacetylnonane-2,8-dione (a photoinitiator according to Example 19 of the present invention). Then, the container was irradiated from above with a Fusion F300 H bulb (Irradiance > 0.25 J/cm²). After the curing process, the content of the container, which contains a cured thick layer on top and uncured liquid material beneath, was removed, and the thickness of the cured top layer was measured by a micrometer screw. Thickness of the cured layer: 800 μm.

INDUSTRIAL APPLICABILITY

The photoinitiator according to this invention is usable for curing of radical curable monomers, oligomers, or
5 polymers.

The photocurable composition of the invention may be used as coatings, printing inks and molded articles. In particularly, the photocurable composition is usable as a coating composition for thick layers of multifunctional
10 acrylic ester groups containing mixtures. That is, the photocurable composition of the invention is characterized in that the composition exhibits excellent photosensitivity even when the coated layer has a thickness of up to 1000 μm .

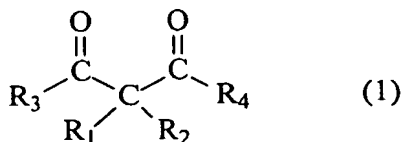
As coatings, the composition can be applied to suitable
15 substrates such as, for example, paper, polyethylene, polypropylene, polyester, polyvinylidene chloride, aluminum, steel or wood and be cured by UV-irradiation in the presence of air or a cover gas.

Among these applications, of particular importance is a
20 coating for durable wood, can coating, or printing ink.

According to this invention, a photoinitiator can be provided, which exhibits excellent photosensitivity, and is usable in thick layer UV curable coatings.

CLAIMS

1. A photoinitiator consisting essentially of a compound having a molecular weight of 1000 or less, and having a chemical structure represented by formula (1),

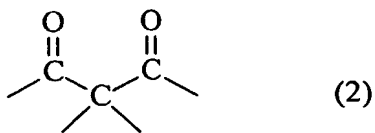


wherein R₃ and R₄ independently denote an alkyl group having a carbon number of 1 to 8, and

R₁ and R₂ independently denote

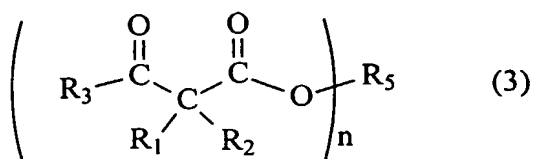
- 1) an electron attracting group,
- 2) an alkyl group having a carbon number of 1 to 8, or
- 3) an alkyl group having a carbon number of 1 to 8, which has an electron attracting group at the β, γ, or δ position with respect to both of the carbonyl groups, wherein the alkyl group 2) is methyl or ethyl group when each of the two substituents is the alkyl group 2), and

weight percentage of a chemical structure element represented by the following formula (2),



- which is expressed in formula (1), based on the total molecular weight of the compound, is within the range of 17% to 54% by mass.

2. A photoinitiator consisting essentially of a compound having a molecular weight of 1000 or less, and having a chemical structure represented by the following formula (3),



5 wherein

R_3 denotes an alkyl group having a carbon number of 1 to 8,

R_5 denotes a mono-, di-, tri-, tetra- or pentavalent aliphatic hydrocarbon group, or an alkyleneoxy group

10 containing aliphatic hydrocarbon group,

n is a natural number of 1 to 5, and

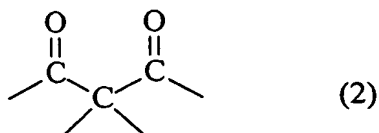
R_1 and R_2 independently denote

1) an electron attracting group,

2) an alkyl group having a carbon number of 1 to 8, or

15 3) an alkyl group having a carbon number of 1 to 8, which has an electron attracting group at the β , γ , or δ position with respect to both of the carbonyl groups, wherein the alkyl group 2) is methyl or ethyl group when each of the two substituents is the alkyl group 2), and

20 weight percentage of a chemical structure element represented by the following formula (2),

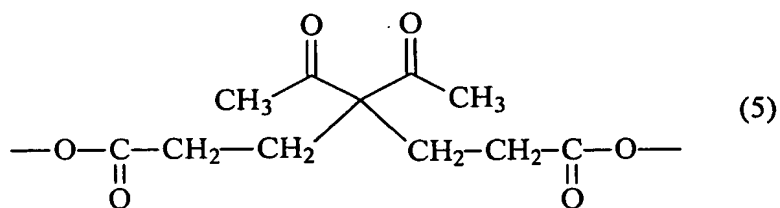


which is expressed in formula (3), based on the total

molecular weight of the compound, is within the range of 17% to 47% by mass.

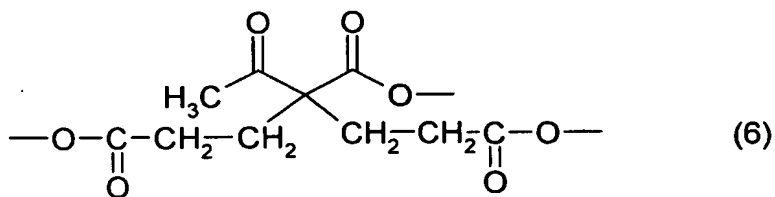
3. A photoinitiator according to claim 1 or 2, wherein the
5 R_1 and R_2 are identical.

4. A photoinitiator according to claim 1, wherein the compound has at least one chemical structure element represented by the following formula (5).



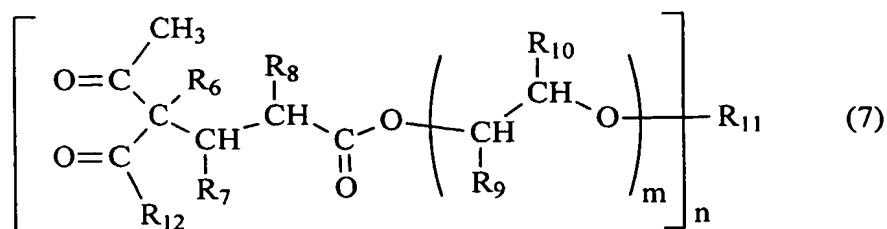
10

5. A photoinitiator according to claim 2, wherein the compound has at least one chemical structure element represented by the following formula (6).



15

6. A novel compound having a chemical structure represented by the following formula (7),



wherein

R₆ denotes an alkyl group having a carbon number of 1 to 8, a C₁₋₄ alkyl carbonyl group, a cyano group, a C₁₋₄ alkyl carbonyl methyl group, a C₁₋₄ alkyl carbonyl ethyl group, a C₁₋₄ alkoxy carbonyl methyl group, a C₁₋₄ alkoxy carbonyl ethyl group, and an alkyl group having a carbon number of 1 to 8 which is substituted by carboxyl group or cyano group,

R₇, R₈, R₉, and R₁₀ independently denote a hydrogen atom, or a methyl group, and at least one of R₉ and R₁₀ is a hydrogen atom,

R₁₁ denotes a di-, tri- or tetra-valent aliphatic hydrocarbon group having a carbon number of 2 to 12,

R₁₂ denotes a methyl group, or an alkoxy group having a carbon number of 1 to 18,

n is a natural number of 2 to 4, and

m is an integer of 0 to 15.

7. A photocurable composition comprising,

- (i) a photoinitiator according to the claim 1 to 7, and
- (ii) a radical curable ethylenic unsaturated compound.

INTERNATIONAL SEARCH REPORT

International Application No

P 03/04282

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C08F2/50 G03F7/031 C07C69/716 C09D4/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C08F C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	GB 2 310 855 A (CIBA SPECIALITY CHEMICALS HOLDING INC.) 10 September 1997 (1997-09-10) cited in the application -----	

☐ Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

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- *O* document referring to an oral disclosure, use, exhibition or other means
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T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

G document member of the same patent family

Date of the actual completion of the international search

30 June 2003

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10/07/2003

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

JP 03/04282

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
GB 2310855	A	10-09-1997	AT 404729 B	25-02-1999
			AT 36297 A	15-06-1998
			BE 1011437 A5	07-09-1999
			BR 9701154 A	15-12-1998
			CA 2198803 A1	04-09-1997
			CH 691970 A5	15-12-2001
			CN 1160718 A , B	01-10-1997
			DE 19708294 A1	11-09-1997
			DK 21797 A	05-09-1997
			ES 2132018 A1	01-08-1999
			FR 2745575 A1	05-09-1997
			IT MI970460 A1	03-09-1998
			JP 10029997 A	03-02-1998
			NL 1005424 C2	22-03-1999
			NL 1005424 A1	05-09-1997
			NO 970962 A	05-09-1997
			RU 2180667 C2	20-03-2002
			SE 9700616 A	04-11-1997
			SG 55281 A1	21-12-1998
			TW 408136 B	11-10-2000
			US 6284813 B1	04-09-2001
			US 6020528 A	01-02-2000
			US 6361925 B1	26-03-2002
			US 2002042022 A1	11-04-2002
			ZA 9701810 A	16-09-1997